CLINICAL PROFILE OF EPILEPSY DURING THE FIRST TWO YEARS OF LIFE

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ABSTRACT:

Objective: This study aims at identifying the different presentations of infantile epilepsy with their E.E.G. and neuro-radiological patterns.

Patient & Methods: Patients included in this study were children suffering from epilepsy starting below 2 years of age. They were attending the outpatient clinic or were from the in-patient departments in Prince Ali - Ben Al - Hussein hospital. The study included 100 children with ages ranging from three month to two years. All patients were subjected to: careful history taking, general and neurological examina-tion including fundus examination, E.E.G. tracing, CT were done for all cases. Febrile convulsions and convulsions due to metabolic disturbances were excluded from the study.

Results: Patients were divided clinically into 2 groups: Group A (symptomatic) group-80 cases (80%): included those who had evident etiology, evident neurological deficit or developmental delay of significant degree prior to onset of seizures while Group B (cryptogenic) group-20 cases (20%). male to female ratio was 1.5: 1. The etiological factors encountered were perinatal asphyxia (55%) while heredofamilial disor-ders were least common 44% of patients had normal clinical examination as well as development while 27 % were mentally handicapped. The generalized tonic-clonic seizure was the commonest in both group while infantile type was the least, EEG was normal in 18 % of cases while CT scan lesion were present in 70% of the patients.

Conclusions: Careful history taking is extremely important for accurate diagnosis of epilepsy. The in-vestigation of choice is the E.E.G. Computerized tomography should be considered an accurate neurodiagnostic test. It provides a good anatomic description of the brain.

KEY WORDS: Infantile epilepsy, presentations, E.E.G.

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INTRODUCTION

Seizure disorder is one of the common child-hood neurological illnesses occurring in 4 - 6 per 1000 children in the general population. The degree and type of central nervous system damage existing at the onset of the seizure is decisive for the final out come, but the cessa-

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tion of the seizures also improves the child's developmental possibilities.²

Determining the type of seizure, and whether it constitutes part of a specific epilepsy syndrome, is important for purposes of work-up, treatment and prognosis. The correct diagnosis of epilepsy leads to an appropriate treatment. It is visible by identifying the etiology or basic disorders underlying the epilepsy by physical and neurological examinations, laboratory tests, including EEGs and neuro-radiological examinations.³

This study aims at identifying the different presentations of infantile epilepsy with their E.E.G. and neuro-radiological patterns to help in their classification and identi-fying whether the condition is cryptogenic or secondary to intracranial pathology. This might be of value in the management of the cases and their outcome.

PATIENTS AND METHODS

Patients included in this study were children suffering from epilepsy starting below 2 years of age. They were attending the outpatient clinic or were from the in-patient departments in Prince Ali – Ben Al - Hussein hospital. The study included 100 children with ages ranging from three month to two years. All patients were subjected to: careful history taking, general and neurological examination including fundus examination, E.E.G. tracing and CT was done in all cases. Febrile convulsions and convulsions due to metabolic disturbances were excluded from the study.

RESULTS

Patients were divided clinically into 2 groups: Group A (symptomatic) 80 cases (80%): included those who had evident etiology, evident neurological deficit or developmental delay of significant degree prior to onset of seizures a Group B (cryptogenic) group-20 cases (20%): this comprised children who had apparently no antecedent etiology, no neurological dis-order and who had normal development until the onset of seizures. the age and six distribution of the patients is presented in Figure-I & II.

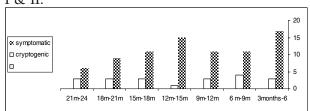


Figure-I: Age distribution of the cases

In all patient male to female ratio was 1.5: 1 male preponderance appear more evident of symptomatic group 3:1 as shown in the Figure-II. The etiological factors encountered were perinatal asphyxia (55%); prema-turity (5%), CNS. infections (15%), head injuries (8%), congenital anomalies of central nervous system (9%) and heredofamilial disor-ders (8%). The generalized tonic-clonic seizure was the commonest in both groups while infantile type was least common (5%) Table-I. EEG was normal in 18 % of cases (abnormal 82.2 % in group A and 56.7 % in group B) as showed in Figure-II.

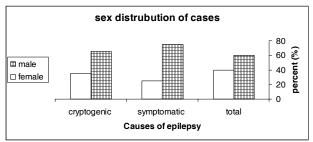


Figure-II: sex distribution of cases

44% of patients had normal clinical examination as well as development while 27 % were mentally handicapped. The head circumference in group A was between 5th and 9th percentile in 27 cases (33.75%) above 95th percentile in 15 % of cases. In group B, it was between 5th percentile and 95th percentile in all cases.

DISCUSSION

Perinatal asphyxia is an important determinant of infant neurological outcome⁴, in our study perinatal asphyxia was the most frequently observed cause accounting for 55 % while other studies have repotted 35%.⁴ It was also reported that seizures are common squeal to sever perinatal asphyxia.^{5,6} In another study it was found that perinatal factors were the most common cause of epilepsy.⁷⁻⁹ Seizure types were partial in 45% of the children and generalized in 55%.

Prematurity was the relevant finding in the history of 5% of the studied cases. Motor and mental disability was present in 75% of them while other studies have reported 33%. They all had generalized seizures. The fits were tonic clonic in 33.3% of cases, myoclonic in 20 %while tonic in others. Brain CT scan showed cortical atrophy in all cases.

CNS infection was an etiological factor for seizures in 15% of cases. This reflects a high incidence of infection among the studied group while earlier studies have reported a lower incidence.¹¹ They found that CNS infection was an etiological factor in 10% of their cases. Some reported an incidence of seizures of 2-8% in a prospective study of 50 infants who recovered from H. in-fluenza meningitis.¹²

Seizures of post traumatic epilepsy may aggravate the brain dysfunction due to primary

trauma and surgical treatment often yields good result and lessens the toxic and adverse effect of antiepileptic drugs.¹³ It was reported that post-traumatic epilepsy represents 4% of the prevalence of the disorder and is one of the sequels which is most difficult to prevent.¹⁴ In our study head injuries formed (8%) which is much lower incidence of early post-traumatic seizures (1.6%) as noted by others.¹⁵ 5% of patients with post-traumatic seizures had normal brain CT –scan as compared to other studies 5.6%.^{16,17}

Table-I: Type of seizure in the studied groups

Seizure type	Group A (%)	Group B (%)	Total (%)
Generalized:	28	6	34
Tonic-clonic	14	3	17
Tonic	7	2	9
Myoclonic	5	2	7
Atonic			
Infantile spase	em 4	2	6
Partial seizure			
Simple	6	5	11
Complex	16		16
Total	80	20	100

Group A : symptomatic Group B : cryptogenic

Neuroimaging examines the relationship between abnormalities of brain function in epilepsy patients (seizures, impaired cognitive function, psychiatric co-morbidity etc.) and focal or more widespread brain pathology. 18 Computed Tomography (CT) is an important tool for neuroimaging as it offers an opportunity to investigate structural lesions as a cause of seizures with little morbidity. 19 CT scan lesion were visible in 70% in the present study, which is comparable to other studies from the subcontinent where it was reported between 50 - 75% but is higher than the prevalence reported in western literature. 20-23

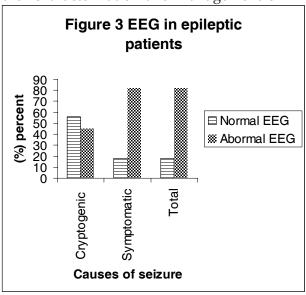
In our study, C.T. abnormalities were present in 70% of group A (it was normal in 30% of cases of group A: 17% of patients had history of birth asphyxia, 5% with t metabolic disorders, 5% had history of meningitis and 3% had history of head trauma. All patients had normal brain CT scan). In patients with generalized seizures, CT abnormalities were detected in 50%. In those with evident neurological

manifestations, CT abnormalities were noted in 78%, compared to 25% abnormal scans with generalized seizures and norm-al neurological findings. In some studies CT findings in children with all varieties of seizures and found an overall abnormality in 33.5%.²⁴

Diagnosis of epilepsy is based upon clinical data. But the electroencephalogram (EEG) remains absolutely necessary for proper management of epileptic patients by the neurologist both in consultation and in hospital practice.²⁵ Normal E.E.G. records were found in (18%) while other studies reported it to be 14%.26 Generalized tonic-clonic were the dominant types of seizures observed in this study (34%) Table-I. It was observed that despite many unusual types of seizures in childhood, the major tonic-clonic convulsions were the com-monest manifestation of childhood epilepsy.²⁷ Myoclonic seizures were observed in 9% of patients in our study while in others it was reported to be 7.5%.²⁷

The work up of 8% cases in the symp-tomatic group led us to the diagnosis of degenerative brain disease. They all showed progressive mental deterioration associated with motor dysfunction affecting the tone, power and reflexes. Macrocephaly was present in two of them. They were offspring of consanguineous marriages of the first degree.

The recent intense focus of attention on further characterization and management of in-



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fantile spasms is due, in part, to the resistant nature of these seizures and the frequently poor cognitive outcome, even when the seizures are controlled.²⁸

Six cases 6% (2% cryptogenic and 4% symptomatic) presented with infantile spasms, usually the onset of their seizures was in the neonatal period. Burst suppression pattern and hypsarrhythmias are age-related EEG patterns that present in infancy and may get modified later on.²⁹ E.E.G. tracings showed hypsarrhythmias and burst suppression which was detected by others who found that perinatal insult was the commonest etiology.^{30,31}

All patients of partial seizures in our study (27%) showed motor seizures while others have reported 48.5 %.⁷ E.E.G. tracing was abnormal in 67% of patients. It was found that the 80% of patients had abnormal neurological examination with partial seizures ³² while in our study it was 75%. Other studies have reported partial seizures and abnormal neurological examination in 95% of patients.¹¹

CONCLUSION

Careful history taking is extremely important for accurate diagnosis of epilepsy. The most important and useful in-vestigation for epilepsy is the E.E.G. Computerized tomography is to be considered an accurate neuro-diagnostic test. It provides a good anatomic descri-ption of the brain. It is especially important in the symptomatic group. Neuroimaging should be considered in any child with a first seizure who does not have an idiopathic form of epilepsy. New techniques such as PET and MRI may provide additional information of the functional disturbances in epilepsy.

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